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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/687,230	10/16/2003	Craig A. Kelly	1895-SPL	8063

7590 03/25/2010
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EXAMINER

KISH, JAMES M

ART UNIT	PAPER NUMBER
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3737

MAIL DATE	DELIVERY MODE
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03/25/2010

PAPER

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UNITED STATES PATENT AND TRADEMARK OFFICE

BEFORE THE BOARD OF PATENT APPEALS
AND INTERFERENCES

Ex parte CRAIG A. KELLY

Appeal 2009-007489
Application 10/687,230
Technology Center 3700

Decided: March 25, 2010

Before LINDA E. HORNER, JENNIFER D. BAHR and
STEVEN D.A. McCARTHY, *Administrative Patent Judges*.

McCARTHY, *Administrative Patent Judge*.

DECISION ON APPEAL

1 The Appellant appeals under 35 U.S.C. § 134 (2002) from the
2 Examiner's decision finally rejecting claims 1-47 under 35 U.S.C. § 103(a)
3 as being unpatentable over Meyer (US 6,308,098 B1, issued Oct. 23, 2001)
4 and Sun (US 6,811,536 B2, issued Nov. 2, 2004). We have jurisdiction
5 under 35 U.S.C. § 6(b) (2002).

1 *Dismissal of the Appeal as to Claims 24-47*

2 The Examiner has entered a new ground of rejection in the
3 Examiner's Answer against claims 24-47 under § 101 as being directed to
4 nonstatutory subject matter. (Ans. 3.) The Examiner properly gave notice
5 of the new ground of rejection. (*Id.*; Ans. 9-10.) The Technology Center
6 Director approved the new ground of rejection. (Ans. 10.) As the Answer
7 indicates (Ans. 9-10), the Appellant was required to respond to the new
8 ground within two months in either of two ways: 1) reopen prosecution, *see*
9 37 CFR § 41.39(b)(1); or 2) maintain the appeal by filing a reply brief as set
10 forth in 37 CFR 41.41, *see* 37 CFR § 41.39(b)(2), "to avoid *sua sponte*
11 dismissal of the appeal as to the claims subject to the new ground of
12 rejection." (Ans. 9.) *See also* 37 CFR § 41.39(b). According to the record
13 before us, the Appellant does not appear to have exercised either option.

14 Accordingly, we DISMISS the appeal as to the claims subject to the
15 new ground of rejection under § 101, namely, claims 24-47. Given that the
16 appeal stands dismissed as to claims 24-47, the sole ground of rejection
17 before us for review is the rejection of claims 1-23 under § 103(a) as being
18 unpatentable over Meyer and Sun.

19 We do not sustain the rejection of claims 1-23. Pursuant to 37 C.F.R.
20 § 41.50(b), we enter a NEW GROUND OF REJECTION against claims 1
21 and 23 under 35 U.S.C. § 101 as being directed to non-statutory subject
22 matter.

23
24 *Reversal of Rejections of Claims 1-23 under § 103(a)*

25 Claims 1, 2, 12, 13 and 23 are independent.

26 Claim 1 recites:

1 1. A non-invasive health monitor device comprising:
2 a processor;
3 a processor readable storage medium;
4 code recorded in the processor readable
5 storage medium to create a first array of data based
6 on discretely recorded time events in which each
7 element of the first array is representative of a time
8 when an event took place;
9 code recorded in the processor readable
10 storage medium to create a second array of data in
11 which each element of the second array is an
12 interval representative of the difference between
13 successive elements of the first array;
14 code recorded in the processor readable
15 storage medium to create a third array of data in
16 which each element of the third array is a delta
17 interval representative of the difference between
18 successive elements of the second array;
19 code recorded in the processor readable
20 storage medium to perform a fast fourier transform
21 (FFT) to obtain power spectrum data
22 representative of the third array; and
23 code recorded in the processor readable
24 storage medium to integrate the power spectrum
25 data over frequency ranges of interest to obtain
26 discrete power values for said frequency ranges of
27 interest.
28

29
30 Claim 23 differs from claim 1 only in that each element of the third
31 array of data is a delta interval representative of the difference between *non-*
32 *successive* rather than successive elements of the second array.

1 Claims 2, 12 and 13 each claim non-invasive health monitor devices.
2 The device of claim 2 includes “code recorded in the processor readable
3 storage medium to create a delta heart period interval array in which each
4 element is a delta heart period interval representative of the difference
5 between successive elements of [a] heart period interval array.” The device
6 of claim 12 includes “code recorded in the processor readable storage
7 medium to create a delta respiration period interval array in which each
8 element is a delta respiration period interval representative of the difference
9 between successive elements of [a] respiration period interval array.” The
10 device of claim 13 includes “code recorded in the processor readable storage
11 medium to create a delta ventricular systole interval array in which each
12 element is a delta ventricular systole interval representative of the difference
13 between successive elements of [a] ventricular systole interval array.”

14 The Examiner provides no clear statement of findings as to the
15 differences between the claimed subject matter and the prior art, namely, the
16 disclosure of the primary reference Meyer. (*See* Ans. 4-5; Final Office
17 Action mailed Sep. 18, 2007 at 3-5; *see also* Ans. 8.) The Examiner fails to
18 articulate any reasoning to explain how the combined teachings of Meyer
19 and Sun would have provided one of ordinary skill in the art reason to
20 *modify* the disclosure of Meyer to provide code to create a “third array of
21 data” as recited in claims 1 and 23; a “delta heart period interval array” of
22 claim 2; a “delta respiration period interval array” as recited in claim 12; or a
23 “delta ventricular systole interval array” as recited in claim 13. (*See id.*)
24 This omission implies that the Examiner finds that Meyer discloses these
25 limitations.

1 The Appellant argues that the “differential analysis step is not an
2 established process or recognized as valuable in analysis of heart rate
3 variability in the prior art . . . Applicant submits that the differential analysis
4 processing step, as recited in all independent claims, is novel and a
5 significant contribution to the measurement and interpretation of heart rate
6 variability.” (Br. 10.) The thrust of this argument is that the combined
7 teachings of Meyer and Sun would have provided one of ordinary skill in the
8 art no reason to program a processor to create the “third array of data” of
9 claims 1 and 23; the “delta heart period interval array” of claim 2; the “delta
10 respiration period interval array” of claim 12; or the “delta ventricular
11 systole array” of claim 13.

12 The Examiner does not identify where Meyer discloses code to create
13 any of these arrays. No such disclosure is evident.

14 Meyer discloses a cardiac pacemaker *100* with the capacity to detect
15 whether a patient is standing or reclining using the morphology of an
16 intracardial physiological sensor signal such as a signal representative of
17 intracardial impedance. (Meyer, col. 2, ll. 6-14.) More specifically,
18 Meyer’s pacemaker *100* includes an impedance signal input stage *101*, an
19 analog-to-digital converter *105*, an impedance signal memory unit *106*, a
20 calibration unit *107* and an arithmetic stage *108*. The elements cooperate to
21 calculate discrete values representing the slopes of an intracardial impedance
22 curve at various time intervals. (Meyer, col. 3, ll. 38-63.) Meyer discloses
23 that:

24 A long-term slope memory *118*, in which
25 (again according to the [“first in-first out”]
26 principle) the slope values from a large number of
27 impedance measurements are stored, is also

1 connected to the output of the arithmetic stage *108*.
2 A fluctuation curve determination stage *119*, in
3 which the curve of the time variability of their
4 fluctuations is determined for all currently stored
5 slope signals, is connected to this. A frequency
6 analyzer *120* for determination of the spectral
7 power density of the fluctuation curve using a
8 linear model of the so-called autoregressive
9 spectral analysis (ARSA) stored in a frequency
10 analysis program memory *121* is connected
11 downstream from this. An integrator stage *122* for
12 integration of the frequency components within the
13 two permanently programmed ranges of 0.05 to
14 0.15 Hz (“LF”) and 0.15 to 0.4 Hz (“HF”) is
15 connected to the output of the frequency analyzer
16 *120*.

17 (Meyer, col. 4, ll. 28-43.)

18 Although the Examiner paraphrases this passage from Meyer in the
19 statement of the grounds of rejection, the Examiner fails to identify how this
20 passage discloses code to create a “third array of data” as recited in claims 1
21 and 23; a “delta heart period interval array” as recited in claim 2; a “delta
22 respiration period interval array” as recited in claim 12; or a “delta
23 ventricular systole interval array” as recited in claim 13. (*See* Ans. 4-5;
24 Final Office Action mailed Sep. 18, 2007 at 3-5; *see also* Ans. 8.) In
25 particular, the Examiner fails to provide reasoning to explain why Meyer’s
26 description of “a fluctuation curve determination stage *119*, in which the
27 curve of the time variability of their fluctuations is determined for all
28 currently stored slope signals, is connected to” the long-term slope memory
29 *118* necessarily discloses these limitations. (*See id.*)

30 Meyer’s statement of the “Objects and Summary of the Invention”
31 describes a process for determining whether a patient is standing or reclining

1 which does not include a differential analysis or the creation of a delta
2 interval array analogous to the “third array of data” of claims 1 and 23; the
3 “delta heart period interval array” of claim 2; the “delta respiration period
4 interval array” of claim 12; or the “delta ventricular systole interval array” of
5 claim 13. (*See Meyer*, col. 2, ll. 35-58.) Meyer’s failure to disclose either a
6 differential analysis or the creation of a delta interval array in the “Objects
7 and Summary of the Invention” indicates that no such disclosure was
8 intended in Meyer’s “Detailed Description of the Invention.”

9 Sun does not remedy the deficiencies of Meyer. Sun teaches
10 determining a heart rate representative of the reciprocals of the time intervals
11 between the peaks of signals of measured blood pressure or
12 electrocardiography. (*Sun*, col. 3, ll. 61-64.) Sun further teaches converting
13 the heart rate to a power spectrum by use of a fourier transform method;
14 separating the power spectrum into high frequency, low frequency and very
15 low frequency variability ranges; and integrating the portion of the power
16 spectrum within each frequency range. (*Sun*, col. 3, l. 64 – col. 4, l. 4.) Sun
17 fails to disclose creating any array which might correspond to the “third
18 array of data” of claims 1 and 23; the “delta heart period interval array” of
19 claim 2; the “delta respiration period interval array” of claim 12; or the
20 “delta ventricular systole interval array” of claim 13.

21 The Examiner has not shown code to create an array corresponding to
22 the “third array of data” of claims 1 and 23; the “delta heart period interval
23 array” of claim 2; the “delta respiration period interval array” of claim 12; or
24 the “delta ventricular systole interval array” of claim 13. The Examiner has
25 not articulated any reasoning which might suggest why one of ordinary skill
26 in the art familiar with the combined teachings of Meyer and Sun might have

1 reason to modify Meyer's pacemaker *100* to include these limitations of
2 claims 1, 2, 12, 13 and 23. We do not sustain the rejections of claims 1, 2,
3 12, 13 and 23 under § 103(a) as being unpatentable over Meyer and Sun.
4 Neither do we sustain the rejections of dependent claims 3-11 and 14-22
5 under § 103(a).

6
7 *New Ground of Rejection of Claims 1 and 23 under § 101*

8 Pursuant to 37 C.F.R. § 41.50(b), we enter a new ground of rejection
9 of claims 1 and 23 under § 101 as being directed to non-statutory subject
10 matter.

11 Subsequent to the mailing of the Examiner's Answer, the Board
12 clarified the scope of statutory subject matter relating to claims reciting
13 machines. A claim to a machine reciting a mathematical algorithm fails to
14 recite statutory subject matter if the claim is not limited to a tangible
15 practical application, in which the mathematical algorithm is applied, that
16 results in a real-world use and not a mere field-of-use label having no
17 significance. *Ex Parte Gutta*, 93 USPQ2d 1025, 1028 (BPAI 2009).

18 Claims 1 and 23 recite machines. Each recites a "non-invasive health
19 monitor device" including at least a processor and a processor readable
20 storage medium.

21 Claims 1 and 23 each recite a mathematical algorithm. Indeed, claims
22 1 and 23 recite no more than the processor, the processor readable storage
23 medium and code for instructing the processor how to perform a
24 mathematical algorithm.

25 Claims 1 and 23 are not limited to *tangible* practical applications.
26 Each recites code to create a first array of data based on discretely recorded

1 time events. Neither claim 1 nor claim 23 limits the manner in which the
2 “data” is obtained. Although both claims require that the “time events” be
3 “discretely record[ed],” neither limits the nature of the “time events” or
4 requires that the time events represent physical events. Each claim also
5 recites code for carrying out four other steps based solely on the first array
6 of data. Nothing recited in claim 1 or claim 23 limits the use of the
7 algorithm to data representing physical objects or real-world process
8 parameters.

9 Claims 1 and 23 do not require that the performance of the algorithm
10 result in a real-world use. The sole result of the algorithm which the code
11 instructs the processor to perform is “discrete power values for . . .
12 frequency ranges of interest.” The discrete power values are not limited to
13 representations of physical objects or real-world process parameters because
14 the power values are derived from “discretely recorded time events” which
15 themselves need not represent physical objects or real-world process
16 parameters. Neither claim requires that any process step be performed in
17 dependence on the magnitude of the discrete power values.

18 Although the preambles of claims 1 and 23 each recite a “non-
19 invasive health monitor device,” this recitation merely identifies an intended
20 use of the claimed device rather than a structural limitation. Neither the
21 body of claim 1 nor the body of claim 23 recites any structure specific to
22 non-invasive health monitoring. Neither claim recites a limitation in the
23 body which relies on the preamble for antecedent basis. Considering the
24 claim language as a whole, the recitation of a “non-invasive health monitor
25 device” in the preamble of each claim merely states a purpose or intended
26 use for the claim and does not positively limit either claim. *See Pitney*

1 *Bowes, Inc. v. Hewlett-Packard Co.*, 182 F.3d 1298, 1305 (Fed. Cir.
2 1999)(discussing circumstances in which preamble recitation do, or do not,
3 limit a claim). Hence, the preamble recitations of claims 1 and 23 are mere
4 field-of-use labels having no significance in limiting claims 1 and 23 to
5 statutory subject matter.

6 Considering the scope of claim 1 and of claim 23 as a whole, neither
7 claim is directed to statutory subject matter.

8
9 DECISION

10 We DISMISS the appeal as to the claims subject to the Examiner's
11 new ground of rejection under § 101, namely, claims 24-47.

12 Upon return of the application to the Examiner, the Examiner should:

13 (1) cancel claims 24-47; and

14 (2) notify the Appellant that the appeal as to the claims
15 subject to the new ground of rejection under § 101 is dismissed
16 and that claims 24-47 are cancelled.

17 See MANUAL OF PATENT EXAMINING PROCEDURE § 1207.03, 8th ed., Rev. 7,
18 Jul. 2008.

19 We REVERSE the decision of the Examiner rejecting claims 1-23.

20 Pursuant to 37 C.F.R. § 41.50(b), we enter a NEW GROUND OF
21 REJECTION of claims 1 and 23 under 35 U.S.C. § 101 as being directed to
22 non-statutory subject matter.

23 Under 37 C.F.R. § 41.50(b) a new ground of rejection has been
24 entered. 37 C.F.R. § 41.50(b) provides that, "[a] new ground of rejection
25 pursuant to this paragraph shall not be considered final for judicial review."

Regarding the new ground of rejection, Appellant must, *WITHIN TWO MONTHS FROM THE DATE OF THE DECISION*, exercise one of the following options with respect to the new ground of rejection, in order to avoid termination of the appeal as to the rejected claims:

(1) *Reopen prosecution*. Submit an appropriate amendment of the claims so rejected or new evidence relating to the claims so rejected, or both, and have the matter reconsidered by the examiner, in which event the proceeding will be remanded to the examiner . . .

(2) *Request rehearing*. Request that the proceeding be reheard under § 41.52 by the Board upon the same record . . .

No time period for taking any subsequent action in connection with this appeal may be extended under 37 C.F.R. § 1.136(a) (2007).

REVERSED; 37 C.F.R. § 41.50(b)

Klh

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